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FILE 'CAPLUS' ENTERED AT 17:25:28 ON 24 MAY 2008

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FILE COVERS 1907 - 24 May 2008 VOL 148 ISS 22

FILE LAST UPDATED: 23 May 2008 (20080523/ED)

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Uploading C:\Program Files\Stnexp\Queries\10517713.str

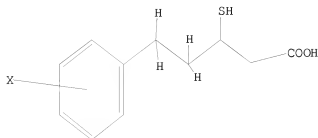
L1 STRUCTURE UPLOADED

=> d

L1 HAS NO ANSWERS

L1 STR

10/923,271



Structure attributes must be viewed using STN Express query preparation.

=> s l1 full

REGISTRY INITIATED

Substance data SEARCH and crossover from CAS REGISTRY in progress...

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FULL SEARCH INITIATED 17:25:52 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 2000 TO ITERATE

100.0% PROCESSED 2000 ITERATIONS

2 ANSWERS

SEARCH TIME: 00.00.01

L2 2 SEA SSS FUL L1

L3 1 L2

=> d ibib abs hitstr

L3 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2008 ACS on SIN

ACCESSION NUMBER: 2003:1006952 CAPLUS

DOCUMENT NUMBER: 140:59517

TITLE: Preparation of 2,5-disubstituted 3-mercaptopentanoic acids as carboxypeptidase U inhibitors

INVENTOR(S): Polla, Magnus

PATENT ASSIGNEE(S): Astrazeneca A.B., Swed.

SOURCE: PCT Int. Appl., 38 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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WO 2003106420	A1	20031224	WO 2003-SE970	20030610

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

CA 2488606	A1	20031224	CA 2003-2488606	20030610
AU 2003241260	A1	20031231	AU 2003-241260	20030610
AU 2003241260	B2	20070426		
BR 2003011384	A	20050315	BR 2003-11384	20030610
EP 1532111	A1	20050525	EP 2003-730987	20030610
EP 1532111	B1	20070808		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
CN 1662504	A	20050831	CN 2003-813840	20030610
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IN 2004DN03722	A	20070302	IN 2004-DN3722	20041124
ZA 2004009955	A	20060830	ZA 2004-9955	20041208
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MX 2004PA12604	A	20050323	MX 2004-PA12604	20041214
HK 1077296	A1	20071109	HK 2005-109105	20051014
PRIORITY APPLN. INFO.:			SE 2002-1837	A 20020614
			WO 2003-SE970	W 20030610

OTHER SOURCE(S): MARPAT 140:59517

AB The title compds. R1(CH2)2CH(SH)CH(CO2H)CH2R2 [I; R1 = (un)substituted Ph, naphthyl, pyridinyl, etc.; R2 = aminopyridinyl, aminothiazolyl, 3-azabicyclo[3.2.1]octyl] which inhibit carboxypeptidase U and thus can be used in the prevention and treatment of diseases where inhibition of carboxypeptidase U is beneficial, were prepared E.g., a 4-step synthesis of 2-[(6-aminopyridin-3-yl)methyl]-5-(1,1'-biphenyl-3-yl)-3-mercaptopentanoic acid (starting from 3-iodo-1,1'-biphenyl), was given. Biol. data was given for 13 exemplified compds. I. In further aspects, the invention relates to pharmaceutical compns. containing at least one compound I.

IT 637300-48-2P 637300-49-3P

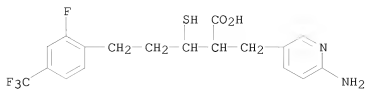
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of 2,5-disubstituted 3-mercaptopentanoic acids as carboxypeptidase U inhibitors)

RN 637300-48-2 CAPLUS

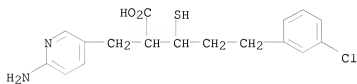
CN 3-Pyridinepropanoic acid, 6-amino- α -[3-[2-fluoro-4-(trifluoromethyl)phenyl]-1-mercaptopropyl]- (CA INDEX NAME)

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RN 637300-49-3 CAPLUS

CN 3-Pyridinepropanoic acid, 6-amino- α -[3-(3-chlorophenyl)-1-mercaptopropyl]- (CA INDEX NAME)



REFERENCE COUNT:

1

THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT